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Term:

L19 and krieg

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File: USPT

Jun 18, 2002

US-PAT-NO: 6406705

DOCUMENT-IDENTIFIER: US 6406705 B1

TITLE: Use of nucleic acids containing unmethylated CpG dinucleotide as an adjuvant

DATE-ISSUED: June 18, 2002

INVENTOR-INFORMATION:

NAME	CITY	STATE	ZIP CODE	COUNTRY
Davis; Heather L.	Ottawa			CA
Schorr; Joachim	Hilden			DE
Krieg; Arthur M.	Iowa City	IA		

US-CL-CURRENT: [424/278.1](#); [424/204.1](#), [424/279.1](#), [424/282.1](#), [536/23.72](#), [930/200](#),
[930/210](#), [930/220](#)

CLAIMS:

We claim:

1. A composition of a synergistic combination of adjuvants, comprising:

an effective amount for inducing a synergistic adjuvant response of a combination of adjuvants, wherein the combination of adjuvants includes at least one oligonucleotide containing at least one unmethylated CpG dinucleotide and at least one non-nucleic acid adjuvant.

2. The composition of claim 1, wherein the non-nucleic acid is an adjuvant that creates a depo effect.

3. The composition of claim 2, wherein the adjuvant that creates a depo effect is selected from the group consisting of alum, emulsion based formulations, mineral oil, non-mineral oil, water-in-oil emulsions, water-in-oil-in-water emulsions, Seppic ISA series of Montanide adjuvants; MF-59; and PROVAX.

4. The composition of claim 1, wherein the non-nucleic acid adjuvant is an immune stimulating adjuvant.

5. The composition of claim 4, wherein the immune stimulating adjuvant is selected from the group consisting of saponins, PCPP polymer; derivatives of lipopolysaccharides, MPL, MDP, t-MDP, OM-174 and Leishmania elongation factor.

6. The composition of claim 1, wherein the non-nucleic acid adjuvant is an adjuvant that creates a depo effect and stimulates the immune system.

7. The composition of claim 6, wherein the adjuvant that creates a depo effect

and stimulates the immune system is selected from the group consisting of ISCOMS, SB-AS2, AS2, SB-AS4, non-ionic block copolymers and SAF.

8. The composition of claim 1, wherein the composition also includes an antigen that is selected from the group consisting of peptides, polypeptides, cells, cell extracts, polysaccharides, polysaccharide conjugates, lipids, glycolipids, carbohydrates, viruses, viral extracts and antigens encoded within nucleic acids.

9. The composition of claim 8, wherein the antigen is derived from an infectious agent selected from the group consisting of a virus, bacterium, fungus and parasite.

10. The composition of claim 8, wherein the antigen is a tumor antigen.

11. The composition of claim 8, wherein the antigen is an allergen.

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